

MIXED HYDROGEN GENERATION MATERIAL

FIELD OF THE INVENTION

[0001] The present invention relates to hydrogen storage compositions, the method of making such hydrogen storage compositions and use thereof for storing hydrogen.

BACKGROUND OF THE INVENTION

[0002] Hydrogen is desirable as a source of energy because it reacts cleanly with air producing water as a by-product. In order to enhance the desirability of hydrogen as a fuel source, particularly for mobile applications, it is desirable to increase the available energy content per unit volume of storage. Presently, this is done by conventional means such as storage under high pressure, at thousands of pounds per square inch, cooling to a liquid state, or absorbing hydrogen into a solid such as a metal hydride. Pressurization and liquification require relatively expensive processing and storage equipment.

[0003] Storing hydrogen in a solid material provides relatively high volumetric hydrogen density and a compact storage medium. Hydrogen stored in a solid is desirable since it can be released or desorbed under appropriate temperature and pressure conditions, thereby providing a controllable source of hydrogen.

[0004] Presently, it is desirable to maximize the hydrogen storage capacity or content released from the material, while minimizing the weight of the material to improve the gravimetric capacity. Further, many current

materials only absorb or desorb hydrogen at very high temperatures and pressures. Thus, it is desirable to find a hydrogen storage material that generates or releases hydrogen at relatively low temperatures and pressures, and which has relatively high gravimetric hydrogen storage density.

[0005] Therefore, in response to the desire for an improved hydrogen storage medium, the present invention provides a method of storing and releasing hydrogen from storage materials, as well as improved hydrogen storage material compositions.

SUMMARY OF THE INVENTION

[0006] In one aspect, the present invention provides a hydrogen storage composition comprising: $M'_xM''_yN_zH_d$ wherein (a) M' is a cation selected from the group consisting of: Li, Ca, Na, Mg, K, Be, and mixtures thereof and x is greater than about 50 and less than about 53; (b) M'' comprises a cation composition comprising a Group 13 element of the Periodic Table and y is greater than about 5 and less than about 34; (c) N is nitrogen and z is greater than about 16 and less than about 45; (d) H is hydrogen and in a fully hydrogenated state, d is greater than about 110 and less than about 177; and (e) wherein M', M'', x, y, z, and d are selected so as to maintain electroneutrality.

[0007] Another aspect of the present invention provides a method of storing hydrogen comprising: reacting a nitride having one or more cations other than hydrogen with a hydride having one or more cations other than hydrogen. The reacting forms a hydrogen storage composition comprising hydrogen,

nitrogen, and at least one of the one or more cations other than hydrogen derived from the nitride and derived from the hydride, respectively.

[0008] Another preferred embodiment of the present invention provides a method of releasing hydrogen comprising: reacting a hydrogen storage nitride composition having one or more cations other than hydrogen with a hydrogen storage hydride composition having one or more cations other than hydrogen, wherein the reacting releases hydrogen and forms one or more byproduct compounds comprising: nitrogen and at least one of the one or more cations other than hydrogen derived from the nitride composition and from the hydride composition, respectively.

[0009] Further areas of applicability of the present invention will become apparent from the detailed description provided hereinafter. It should be understood that the detailed description and specific examples, while indicating the preferred embodiment of the invention, are intended for purposes of illustration only and are not intended to limit the scope of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

[0010] The present invention will become more fully understood from the detailed description and the accompanying drawings, wherein:

[0011] Figure 1 shows a high-resolution x-ray diffraction pattern of a new quaternary hydrogen storage compound;

[0012] Figure 2 shows x-ray diffraction patterns of a mixture of reactants milled according to the present invention compared with the new hydrogen storage compound;

[0013] Figure 3 shows x-ray diffraction patterns of starting reactants compared with the new hydrogen storage compound;

[0014] Figure 4 shows thermogravimetric analysis (TGA) of the weight loss of the new hydrogen storage compound versus time as temperature is increased to 250°C and then held constant, combined with residual gas analysis (RGA) data obtained with a mass spectrometer monitoring the exhaust gas from the TGA; and

[0015] Figure 5 shows another TGA of the new hydrogen storage compound, where heat is applied at a constant rate of 5°C per minute while an RGA monitors the evolved gas.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0016] The following description of the preferred embodiment(s) is merely exemplary in nature and is in no way intended to limit the invention, its application, or uses.

[0017] In one aspect, the present invention provides a method of storing and releasing hydrogen. In one feature, a hydrogen storage material is formed by combining a nitride and a hydride, each of which are solids. The hydride is preferably represented by the general formula $MI^a(MIIC_{b,a})_a$, where MI represents a first cationic species other than hydrogen, MII represents a second cationic species other than hydrogen, a represents an average valence state of MI and $\left(\frac{a \times b}{1+a}\right)$ represents an atomic ratio of hydrogen to cationic species (i.e., MI and MII) in the hydride compound. The nitride is preferably represented by the general formula $MIII^f(NH_e)_g^{-c}$, where MIII represents a cationic species other than hydrogen, N represents nitrogen, H represents hydrogen, f represents an average valence state of MIII, $c = (3 - e)$, $g = \frac{f}{c}$ and $\left(\frac{e \times g}{1+g}\right)$ represents the atomic ratio of hydrogen to cationic species (i.e., MIII, N) in the nitride compound. Thus, MI, MII, and MIII in both the hydride and the nitride respectively represent cationic species or a mixture of cationic species other than hydrogen.

[0018] A novel hydrogen storage composition material is formed as a stable intermediate (SI) in the reaction of the nitride with the hydride, as

described above. As used herein, the term "composition" refers broadly to a substance containing at least the preferred chemical compound, but which may also comprise additional substances or compounds, including impurities. The term "material" also broadly refers to matter containing the preferred compound or composition. The formation of such a SI compound is dependent upon the individual chemical characteristics of the metal hydride and the nitride selected, and thus is most thermodynamically favored for certain preferred reactions. The SI hydrogen storage material is preferably in a solid phase form, and most preferably in a single solid phase form. The SI hydrogen storage composition preferably comprises hydrogen, nitrogen, and at least one of the one or more cations other than hydrogen derived from the nitride and derived from the hydride, respectively. The SI hydrogen storage composition further undergoes a decomposition reaction where the stored hydrogen is released. The products of this decomposition reaction are hydrogen and one or more byproduct compounds comprising nitrogen, and the one or more cations other than hydrogen derived from both the nitride compound and the hydride compound, respectively.

[0019] Thus, the present invention provides a method of storing hydrogen in a quaternary SI hydrogen storage composition. The reaction between the nitride and hydride compounds, described above, forms the stable quaternary intermediate (the novel hydrogen storage compound). Subsequent to the formation of the SI, hydrogen may be stored at ambient conditions in a stable form. When the release of hydrogen is desired, heat and/or pressure are

applied to facilitate a decomposition reaction, where hydrogen gas is released from the quaternary SI hydrogen storage compound, and one or more decomposition byproducts are formed.

[0020] In another aspect, the present invention provides a method of releasing and generating hydrogen by reacting a nitride having one or more cations other than hydrogen with a hydride, also comprising one or more cations besides hydrogen. The nitride and hydride compounds react to release and form hydrogen and one or more byproduct compounds. In such methods of the present invention, the nitride and hydride react to directly produce hydrogen via reaction, rather than to form a stable intermediate. Whether the SI forms is related to the thermodynamics of each reaction, and appears not to form in some reactions, either due to the instability of any intermediate that may form, or because the reaction does not appear to produce any intermediate, but rather directly proceeds to the final reaction products (i.e., hydrogen and the one or more substantially dehydrogenated byproduct compounds).

[0021] Thus, in certain preferred embodiments, the present invention provides two distinct physical states, one where hydrogen is “stored” and another subsequent to hydrogen release. Where the starting reactants react without forming an SI, the hydrogenated storage state corresponds to the reactants (i.e., because a stable hydrogenated intermediate is not formed), and the byproduct compound(s) correspond to the dehydrogenated state.

[0022] It should be understood that in the present invention MI, MII, MIII of the nitride and hydride compounds, previously described, each represent

a cationic species or mixture of cationic species other than hydrogen. Examples are metal cations, non-metal cations such as boron, and non-metal cations which are organic such as CH₃. Elements that form preferred nitrides, hydrides, and mixtures of cations in the type of compounds of the present invention are as follows. Preferred cationic species generally comprise: aluminum (Al), arsenic (As), boron (B), barium (Ba), beryllium (Be), calcium (Ca), cadmium (Cd), cerium (Ce), cesium (Cs), copper (Cu), europium (Eu), iron (Fe), gallium (Ga), gadolinium (Gd), germanium (Ge), hafnium (Hf), mercury (Hg), indium (In), potassium (K), lanthanum (La), lithium (Li), magnesium (Mg), manganese (Mn), sodium (Na), neodymium (Nd), nickel (Ni), lead (Pb), praseodymium (Pr), rubidium (Rb), antimony (Sb), scandium (Sc), selenium (Se), silicon (Si), samarium (Sm), tin (Sn), strontium (Sr), thorium (Th), titanium (Ti), thallium (Tl), tungsten (W), yttrium (Y), ytterbium (Yb), zinc (Zn), and zirconium (Zr), and organic cations including (CH₃) methyl groups.

[0023] Nitride compounds, as used herein, include nitrogen-containing compounds having one or more cationic species, as described above, and hydrogen. The term "nitride" broadly includes compounds comprising amides (NH₂ group), imides or nitrenes (NH group), and azides (N₃ group). For nitride compounds, preferred cationic species (M_{III}) comprise Al, B, Ca, Li, Na, K, Be, Sr and Mg. Particularly preferred nitride compounds according to the present invention comprise the following non-limiting examples, lithium amide (LiNH₂), sodium amide (NaNH₂), lithium nitride (Li₃N), borozane, also known as borane-

ammonia complex, (BNH_6), lithium azide (LiN_3), magnesium amide ($\text{Mg}(\text{NH}_2)_2$), magnesium imide (MgNH), and mixtures thereof.

[0024] Metal hydride compounds, as used herein, include those compounds having one or more cations other than hydrogen, and may comprise complex metal hydrides, which include two or more distinct cations other than hydrogen (i.e. M_I and M_{II} are distinct cationic species), as previously described. Particularly preferred M_I and M_{II} cations for hydrides comprise cations selected from the group: Al, B, Ca, Li, Na, and Mg. In certain preferred embodiments, it is preferred that M_I and M_{II} are different species, forming the complex metal hydride. In certain embodiments, the metal hydride compound may have one or more cations that are selected from a single cationic species (i.e. M_I and M_{II} are the same cationic species). Preferred metal hydrides according to the present invention comprise the following non-limiting examples, lithium hydride (LiH), lithium aluminum hydride (LiAlH_4), sodium borohydride (NaBH_4), lithium borohydride (LiBH_4), magnesium borohydride ($\text{Mg}(\text{BH}_4)_2$) and sodium aluminum hydride (NaAlH_4).

[0025] It should be noted that M_I, M_{II}, and M_{III} are independently selected in both the nitride and metal hydride compounds, and each may be different, or any two or more may be the same, cationic species. In certain preferred embodiments according to the present invention, M_I and M_{III} are the same cationic species in both the nitride and the metal hydride, however, it is within the scope of the present invention to have distinct cationic species for M_I of the nitride and the M_{III} of the metal hydride. Further, M_{II} may be the same as

MI in the metal hydride, as previously discussed, creating a metal hydride with a single cationic species.

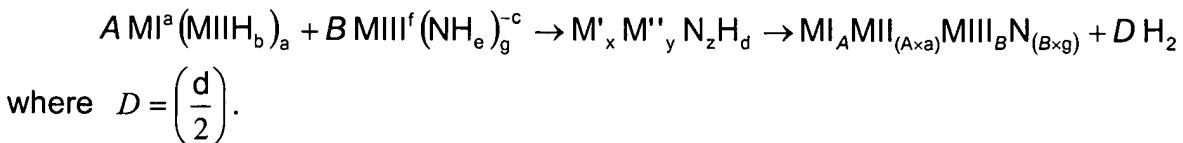
[0026] Thus, according to one preferred embodiment of the present invention, the general reaction for releasing hydrogen while reacting a nitride and metal hydride proceeds according to the following mechanism:



$$c = (3 - e), D = \frac{(A \times a \times b) + (B \times e \times g)}{2}, \text{ and } g = \frac{f}{c}, \text{ and } a, b, c, e, f, g, A \text{ and } B,$$

are selected so as to maintain electroneutrality. It should be noted that the byproduct compound $\text{MI}_A \text{MII}_{(A \times a)} \text{MIII}_B \text{N}_{(B \times g)}$ may thermodynamically favor decomposing into further smaller and/or distinct byproduct compounds. These further byproducts are formed of the same general constituents as the primary byproduct, but they have different valence states, atomic ratios, or stoichiometry, depending on the cationic species involved, as recognized by one of skill in the art. Such additional distinct byproduct compounds may include metal hydrides, which may slightly detract from the total amount of hydrogen generated designed as $D\text{H}_2$.

[0027] As previously discussed, in certain preferred embodiments a stable intermediate hydrogen storage compound is formed, which is expressed by the following reaction, proceeding by a similar reaction mechanism to the general reaction shown above:



[0028] Although not wishing to be limited to any particular theory, a novel solid quaternary intermediate compound is known to occur where the metal hydride has one or more M' cations selected as Li, and generally believed to occur where M' is selected from the group consisting of: Li, Ca, Na, Mg, K, Be, and mixtures thereof, and where M" comprises a cation compound comprising a Group 13 element from the IUPAC Periodic Table. Where the novel SI hydrogen storage composition is formed, such a composition is represented by the general formula $M'_xM''_yN_zH_d$, where N is nitrogen and H is hydrogen. As can be observed in the mechanism above, such a compound undergoes a decomposition reaction mechanism, to form a dehydrogenated state where one or more decomposition byproducts, represented generally by the formula $MI_AMII_{(Ax)}MIII_BN_{(Bx)}$ are formed in addition to a hydrogen product, represented by the general formula, DH_2 . It should be noted that the M' and M" are formed from the MI, MII, and MIII cations present in the reactants, and may comprise one or more cations, including mixtures thereof. Preferably, the MI and MIII cations are the same, and form the M'. Further, in preferred embodiments, x is greater than about 50 and less than about 53; y is greater than about 5 and less than about 34; z is greater than about 16 and less than about 45; d is greater than about 110 and less than about 177; it is preferred that $x+y+z = 100$; and M', M", x, y, z, and d are selected so as to maintain electroneutrality of the compound.

[0029] Examples of preferred reactions according to the present invention which form a SI hydrogen storage composition comprise:

1) $\text{LiBH}_4 + 2 \text{ LiNH}_2 \rightarrow \text{Li}_3\text{BN}_2\text{H}_8 \rightarrow \text{Li}_3\text{BN}_2 + 4 \text{ H}_2$, which generates a theoretical 11.8 wt% hydrogen of the starting reactants, and a new stable intermediate hydrogen storage compound, $\text{Li}_3\text{BN}_2\text{H}_8$.

2) $\text{LiAlH}_4 + 2 \text{ LiNH}_2 \rightarrow \text{Li}_3\text{AIN}_2\text{H}_8 \rightarrow \text{Li}_3\text{AIN}_2 + 4 \text{ H}_2$, which generates a theoretical 9.5 wt% hydrogen of the starting reactants, and according to predicted thermodynamics favors forming a stable intermediate compound, $\text{Li}_3\text{AIN}_2\text{H}_8$.

[0030] Other non-limiting examples of alternate preferred embodiments according to the present invention where hydrogen generation occurs, and where a stable SI hydrogen storage composition although possible, is less favored to form, prior to the hydride generating reaction based on predicted thermodynamics, include the following exemplary reactions:

3) $\text{NaBH}_4 + 2 \text{ NaNH}_2 \rightarrow \text{Na}_3\text{BN}_2 + 4 \text{ H}_2$ which generates a theoretical 6.9 wt% hydrogen of the starting reactants, where a predicted intermediate compound is $\text{Na}_3\text{BN}_2\text{H}_8$.

4) $\text{Mg}(\text{BH}_4)_2 + 6 \text{ MgNH} \rightarrow 2 \text{ Mg}_3\text{BN}_3 + \text{Mg} + 7 \text{ H}_2$ which generates a theoretical 4.8 wt% of the starting reactants and forms two by-products of the cationic species: magnesium boroazide Mg_3BN_3 and magnesium Mg.

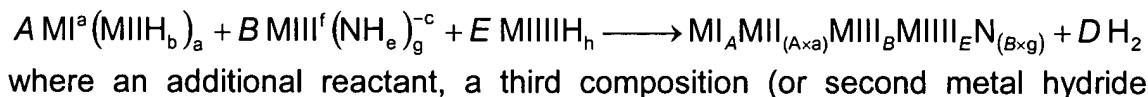
5) $\text{Mg}(\text{BH}_4)_2 + 6 \text{ MgNH} \rightarrow 2 \text{ Mg}_3\text{BN}_3 + \text{MgH}_2 + 6 \text{ H}_2$ which generates a theoretical 4.1 wt% hydrogen of the starting reactants and forms two by-products of the cationic species, magnesium boroazide Mg_3BN_3 and magnesium hydride MgH_2 .

6) $\text{Mg}(\text{BH}_4)_2 + 2 \text{ Mg}(\text{NH}_2)_2 \rightarrow \text{Mg}_3\text{B}_2\text{N}_4 + 8 \text{ H}_2$ which generates a

theoretical 9.6 wt% hydrogen of the starting reactants.

7) $\text{Mg}(\text{BH}_4)_2 + 4 \text{ MgNH} \rightarrow \text{Mg}_3\text{B}_2\text{N}_4 + 2 \text{ MgH}_2 + 4 \text{ H}_2$ which generates a theoretical 3.8 wt% hydrogen of the starting reactants.

[0031] In alternate preferred embodiments according to the present invention, the hydrogen generation reaction proceeds according to the reaction mechanism:



(MIIIIH_h)) is present, and $c = (3 - e)$, $D = \frac{(A \times a \times b) + (B \times e \times g) + (E \times h)}{2}$, and $g = \frac{f}{c}$

and a, b, c, e, f, g, h, A, B, D, and E are selected so as to maintain electroneutrality. Examples of exemplary preferred reactions according to the above mechanism having a third reactant composition include:

8) $\text{Mg}(\text{BH}_4)_2 + 3 \text{ Mg}(\text{NH}_2)_2 + 2 \text{ Mg} \rightarrow 2 \text{ Mg}_3\text{BN}_3 + 10 \text{ H}_2$ which generates a theoretical 7.4 wt% hydrogen of the starting reactants with additional magnesium added as a reactant, where an intermediate although possible is less favored by predicted thermodynamics to form, and a byproduct consisting mainly or solely of magnesium boroazide Mg₃BN₃ does form.

9) $\text{Mg}(\text{BH}_4)_2 + 3 \text{ Mg}(\text{NH}_2)_2 + 2 \text{ MgH}_2 \rightarrow 2 \text{ Mg}_3\text{BN}_3 + 12 \text{ H}_2$ which generates a theoretical 8.7 wt% hydrogen of the starting reactants (where an additional reactant is present, magnesium hydride MgH₂), and a byproduct consisting mainly of magnesium boroazide Mg₃BN₃ is favored to form.

[0032] Preferred conditions for reaction of the nitride compound with the metal hydride compound vary with respect to preferred temperature and

pressure conditions for each independent reaction. However, it is preferred that the reaction is carried out as a solid state reaction, in a non-oxidizing atmosphere, essentially in the absence of oxygen, preferably in an inert atmosphere, such as under nitrogen or argon. Further, as will be discussed in more detail below, it is preferred that the solid nitride and metal hydride compounds are reduced in particle size from their starting size. In the case of the nitride, an average particle diameter size of less than about 3 μm is preferred, and for the metal hydride, an average particle diameter size of less than 25 μm (microns) and most preferably to less than 15 μm is desirable. The reduction of particle size may occur prior to conducting the reaction or concurrently to conducting the reaction between the compounds. In certain preferred embodiments, the reaction is carried out at ambient pressure at a temperature of about 85°C or higher. However, as recognized by one of skill in the art, such temperatures and pressures are highly dependent on the reaction kinetics for each individual reaction.

[0033] With reference to the embodiment where the SI hydrogen storage composition (represented by the general formula $\text{M}'_x \text{M}''_y \text{N}_z \text{H}_d$) is formed, it is preferred in certain embodiments, that an alkali metal hydride is reacted with an alkali nitride. One preferred example is where the lithium is the alkali metal cationic species. The formula unit (and corresponding atomic ratios) of the intermediate compound is best expressed by $\text{Li}_x \text{B}_y \text{N}_z \text{H}_d$, where preferred ranges for x are greater than about 50 and less than about 53; preferred ranges for y are greater than about 5 and less than about 34, preferred ranges for z are

greater than about 16 and less than about 45, and preferred ranges for *d* are greater than about 110 and less than about 177. Further, *x*, *y*, *z*, and *d* are selected so as to maintain the electroneutrality of the hydrogen storage intermediate compound. The SI hydrogen storage compound may be represented by the simplified general formula $\text{Li}_q\text{B}_r\text{N}_s\text{H}_t$, where the atomic ratios may be expressed by the following relationships: *q/r* is about 3; *s/r* is about 2; and *t/r* is about 8. Thus, the average atomic ratio of one preferred SI can be expressed by the nominal general formula $\text{Li}_3\text{BN}_2\text{H}_8$. In certain preferred embodiments of the present invention, the compounds which form the lithium SI compound are a lithium hydride reacted with a lithium nitride. Such lithium hydrides may include, for example, LiAlH_4 , LiH , LiBH_4 . Lithium nitrides may include and LiNH_2 , Li_3N , BNH_6 , and LiN_3 .

[0034] In one preferred reaction, the reactants for the reaction forming the $\text{Li}_x\text{B}_y\text{N}_z\text{H}_d$ hydrogen storage composition are lithium amide compound and lithium borohydride compound. The preferred stoichiometry in the following reaction $A \text{ LiBH}_4 + B \text{ LiNH}_2 \rightarrow \text{Li}_x\text{B}_y\text{N}_z\text{H}_d$ is preferably a stoichiometric ratio of nitride to metal hydride (A:B) from between about 0.5 to about 3. Particularly preferred stoichiometric ratios of A:B are where *A* is about 1 and *B* is between about 2 to about 2.25, which corresponds to an *x* of about 50, a *y* of about 15 to about 17, a *z* of about 33 to about 35, and an *d* of about 130 to about 134. For this reaction, the temperature of formation at ambient pressure is from about 85°C to about 95°C.

[0035] After the novel Si hydrogen storage composition is formed, it is a hydrogenated and stable material at ambient conditions. When release of the hydrogen is desired, the composition is heated to a temperature of about 170°C at ambient pressure. The melting point of the hydrogen storage composition is about 210°C at ambient pressure. Hydrogen release has been observed to occur much more rapidly when the Si hydrogen storage composition is in a liquid state, versus a solid or partially solid state, and thus according to the present invention, it is preferred that the compound is heated to above the melting point of the composition to rapidly release the hydrogen gas.

[0036] A first experiment was conducted according to a method of making a hydrogen storage compound according to a first preferred embodiment of the present invention, where a mixture of LiBH₄ and LiNH₂ of molar ratio 1:2 react according to the above described chemical reaction formula. The LiBH₄ is commercially available from Lancaster Synthesis, Inc. of Windham, New Hampshire (and is specified to be ≥95% purity) and the LiNH₂ is commercially available from Aldrich Chemical Co. of St. Louis, Missouri (also specified to be ≥95% purity). The starting powders are mixed in the molar ratio 1 LiBH₄ : 2 LiNH₂ and sealed into a hardened steel ball mill jar while inside an argon (Ar) inert atmosphere glove box. One large and two small steel balls are placed in the jar with the powder. The material is then high-energy ball milled for at least five hours using a SPEX 8000 mixer-mill. The resulting powder is Li₃BN₂H₈. After 16 hours of milling, the average particle size diameter of the

compound(s) remaining in the mill were measured to be approximately 6 μm to about 11 μm .

[0037] In a second experiment according to an alternate preferred embodiment of the present invention, a method of making a hydrogen storage compound is provided, the ratio of reactants to one another is the same as described in the embodiment above. In the second experiment, the LiBH₄ is sealed into a steel ball mill jar along with one large and two small steel balls while inside an Ar inert gas glove box. The powder is then milled for 10 minutes using a SPEX 8000 mixer mill in order to reduce the average powder particle diameter size to about 6 μm to about 11 μm . The same process is used to separately mill LiNH₂ for 10 minutes for the same purpose, where the resulting particle size is about 1.5 μm to about 3 μm .

[0038] As observed by X-ray diffraction, no change in composition or crystal structure of either reactant occurs after milling the reactant by itself. The two pre-milled powders are then weighed out in the molar ratio 1 LiBH₄ : 2 LiNH₂ and once more sealed into a ball mill jar under Ar gas, but this time without including any balls. The jar is then agitated in the SPEX 8000 mixer-mill for 10 minutes. In the absence of milling balls, the only effect is to thoroughly homogenize the mixture of powders. X-ray diffraction performed after the mixing confirms that no transformation has occurred yet. The mixed powder is then heated above about 90°C for a short time (less than 10 minutes) to convert it via reaction to Li₃BN₂H₈ powder.

[0039] Figure 1 shows a high-resolution x-ray diffraction pattern (the x-ray “fingerprint”) of the new quaternary hydride, or Si hydrogen storage composition, referred to generally as $\text{Li}_3\text{BN}_2\text{H}_8$ formed in the first experiment by ball milling for 16 hours. The vertical lines indicate the angles 2θ of the experimentally observed diffraction lines. There is one extraneous peak at about $2\theta = 21.3^\circ$ due to a plastic film that was placed over the sample surface to prevent the sample from reacting with oxygen in the air. Evidence that this pattern represents a single unique phase comes from two observations. First, the same pattern of lines (both 2θ values and intensities) occurs for samples produced under a variety of experimental conditions. Second, preliminary analysis of the peak positions indicates that a single model for the crystal symmetry can explain all of the observed lines.

[0040] Figure 2 illustrates the effect of ball milling as performed in the first experiment on the x-ray diffraction pattern of the mixture $\text{LiBH}_4 + 2 \text{ LiNH}_2$. The lowest diffraction pattern, labeled “0” milling time, is for the starting mixture, and is comprised of lines characteristic of LiBH_4 (represented by inverted triangles) and LiNH_2 (represented by circles). A trace quantity of Li_2O (represented by diamonds) is also present as an impurity. The center patterns show the evolution of the diffraction pattern with ball milling time, from 10 minutes of milling up to 300 minutes. Patterns for 80, 120, and 200 minutes have been omitted for clarity; they differ in only small details from the 160 minutes pattern. For comparison, the pattern obtained after much longer milling time, 960 minutes, is also shown, but it does not differ from the 300 minutes

pattern, indicating that the transformation is complete after 300 minutes. The squares, together with the vertical gray lines, indicate the positions of the diffraction lines for $\text{Li}_3\text{BN}_2\text{H}_8$ obtained from the x-ray diffraction analysis of Figure 1.

[0041] As milling proceeds, the diffraction pattern of LiBH_4 weakens (best observed from the quartet of lines between 24° and 27°), as does the pattern for LiNH_2 (seen from the main peak at 30.5°). Concurrently the pattern for $\text{Li}_3\text{BN}_2\text{H}_8$ emerges. As the transformation proceeds there are small changes in the positions and intensities of the $\text{Li}_3\text{BN}_2\text{H}_8$ lines (for example, the main peak at 28.7°) indicating that the details of the underlying crystal structure also evolve with milling. The Li_2O content is essentially unchanged, but the $\text{Li}_3\text{BN}_2\text{H}_8$ lines obscure its diffraction lines. In summary, x-ray diffraction shows that ball milling of LiBH_4 and LiNH_2 (in a 1:2 molar ratio) induces a transformation to the new hydrogen storage compound, which is a quaternary hydride phase $\text{Li}_3\text{BN}_2\text{H}_8$.

[0042] Figure 3 shows the effect of heating on the sample mixture from the second experiment comprising LiBH_4 and 2 LiNH_2 . The mixture transforms to $\text{Li}_3\text{BN}_2\text{H}_8$ when it reaches a temperature of about 90°C . Shown are characteristic diffraction patterns obtained below (71°C) and above (128°C) the transformation temperature. The original sample from the second experiment was a mixture of premilled LiBH_4 and LiNH_2 . First the LiBH_4 and the LiNH_2 were prepared in accordance with the second embodiment described above, where each reactant was milled separately for 10 minutes to reduce the particle size, and then mixed without grinding media for an additional 10 minutes to

homogenize the mixture. The mixture was then stored in an Ar inert atmosphere glove box at room temperature for 10 days. The mixture formed the sample for the heat treatment experiment used for the results shown in Figure 3, and contains a substantial quantity of $\text{Li}_3\text{BN}_2\text{H}_8$ in addition to the original LiBH_4 and LiNH_2 , as shown by the 71°C diffraction pattern in Figure 3 (the room temperature diffraction pattern is essentially identical to the 71°C pattern). Although not limiting to the mechanism by which the present invention operates, this implies that the mixed reactant powders ($\text{LiBH}_4 + 2 \text{ LiNH}_2$) are slowly converting to $\text{Li}_3\text{BN}_2\text{H}_8$ at room temperature. When heated to temperatures above 90°C , the sample converts entirely to the $\text{Li}_3\text{BN}_2\text{H}_8$ phase, as shown by the 128°C diffraction pattern. From the temperature evolution of the diffraction pattern, the transformation occurs between about 85°C to about 95°C , and the duration for complete transformation was less than 10 min.

[0043] Figure 4 shows a thermogravimetric analyzer (TGA) analysis with a coinciding residual gas analysis (RGA) obtained with a mass spectrometer monitoring the exhaust gas from the TGA. The upper panel of Figure 4 shows hydrogen release by weight loss as a function of time for a sample of $\text{Li}_3\text{BN}_2\text{H}_8$. The sample used to develop Figure 4 is from a third experiment, where the sample of a stable quaternary hydride hydrogen storage compound is formed by mixing $\text{LiBH}_4:\text{LiNH}_2$ in the ratio 1:2 (having 5 wt% TiCl_3 as an inert impurity). The mixed powder was ball milled using a SPEX 8000 mixer-mill for 10 minutes. X-ray diffraction showed the as-milled powder to consist mostly of the $\text{Li}_3\text{BN}_2\text{H}_8$ phase, with some retained LiBH_4 and LiNH_2 .

Other experiments have shown that the remaining LiBH₄ and LiNH₂ will convert to Li₃BN₂H₈ upon heating above 90°C. Although TiCl₃ sometimes acts as a catalyst for some hydrogen storage materials, such as NaAlH₄, the TiCl₃ appears to be completely inert in the LiBH₄-LiNH₂ system.

[0044] The sample was heated in steps of 50°C to 250°C (represented by the dash-dot line). At 250°C the sample lost weight continuously over time until the total weight loss reached 13.3 wt%, at which point the weight loss stopped abruptly. The weight loss is slightly larger than the theoretical hydrogen content of the sample (11.3 wt%). Although not wishing to be bound by any particular theory, the greater weight loss may be attributed to decomposition of one or more of the impurity phases present in about 5% levels in the starting materials, or possibly due to a slight loss of nitrogen due to producing a small amount of ammonia NH₃, or both.

[0045] As shown in the lower panel of Figure 4, the RGA signals are depicted for mass 2 (H₂), mass 17 (NH₃), and mass 16 (a “crack” of NH₃ corresponding to NH₂⁻). The estimated contribution to mass 17 due to a crack of H₂O corresponding to OH⁻ has been subtracted from the mass 17 signal. Also the mass 17 signal has been shifted upward (multiplied by 2 on a log scale) purely for display purposes to separate it from the mass 16 signal. The signal near time t=0 represents the background signal from the RGA in the absence of these mass species in the sampled gas. Starting at 100°C, heating is accompanied by bursts of evolved gas identified as a mixture of H₂ and NH₃. At 250°C almost all of the evolved gas is H₂. The abrupt end of weight loss after

900 minutes is strongly correlated with the sudden shutoff of the H₂ signal in the RGA, clearly demonstrating that the bulk of the evolved gas is hydrogen.

[0046] Figure 5 shows an alternative TGA measurement in which a sample of Li₃BN₂H₈ (without the additive TiCl₃) is heated to 350°C at a constant rate of 5°C per minute. The RGA was used to monitor the composition of the exhaust gas and identify the evolved gas species, as shown in the lower panel of Figure 5, which corresponds to the time frame of the upper panel. The correlation between the TGA and RGA again demonstrates that the sample simultaneously evolved both H₂ and NH₃ gas. The results indicate that rapid weight loss starts at temperatures of around 275°C, but this is an upper limit since there is a time delay of up to several minutes between the actual emission of the gas by the sample inside the TGA, and its appearance in the exhaust gas stream where it is detected by the RGA. Semiquantitative analysis of the RGA signals estimates that the total weight loss due to H₂ is about 12.2 wt% (in reasonable agreement with the theoretical hydrogen content of 11.9 wt%) and about 1.7 wt% NH₃.

[0047] Although the reversibility of the reaction does not appear to presently occur at a sufficient rate for suitable temperature and pressure conditions desirable for a commercial embodiment, incorporating a catalyst is one known method to both reduce the hydrogen release temperature and facilitate reabsorption of hydrogen in other prior art hydrogen storage materials. Thus, the present invention contemplates employing such a catalyst, as known to one of skill in the art, to facilitate reversibility at desirable conditions and rates.

Catalysts that may be useful with the present invention, include, for example, the following non-limiting list: Fe, Ni, Co, Pt, Pd, Sr, and compounds and mixtures thereof.

[0048] Thus, the hydrogen storage materials according to the present invention provide a novel stable solid phase hydrogen storage composition material, which is especially advantageous in fuel cell applications. The reaction to generate hydrogen is readily controlled by temperature and pressure, and the solid phase is capable of storing hydrogen for prolonged periods at moderate conditions. Further, the methods of the present invention provide a method of releasing hydrogen from solid materials, which can be controlled both by limiting when the reactants are mixed, or under which conditions (e.g. temperature, pressure, and reactant particle size) the reactants are mixed, to provide a controlled hydrogen release from solid phase hydrogen storage materials.

[0049] The description of the invention is merely exemplary in nature and, thus, variations that do not depart from the gist of the invention are intended to be within the scope of the invention. Such variations are not to be regarded as a departure from the spirit and scope of the invention.